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Frank C. Eisenschenk, Ph.D., Patent Attorney

REQUEST FOR CERTIFICATE OF
CORRECTION UNDER 37 CFR 1.322
AND UNDER 37 CFR 1.323
Docket No. ARS.124

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicants : Amanda Proudfoot, Jeffrey Shaw, Zoe Johnson
Issued : June 22, 2010
Patent No. : 7,740,833
Conf. No. : 9024
For : Therapeutic Uses of Chemokine Variants

Mail Stop Certificate of Corrections Branch
Commissioner for Patents
P.O. Box 1450
Alexandria, VA 22313-1450

REQUEST FOR CERTIFICATE OF CORRECTION
UNDER 37 CFR 1.322 (OFFICE MISTAKE) AND
UNDER 37 CFR 1.323 (APPLICANT MISTAKE)

Sir:

A Certificate of Correction for the above-identified patent has been prepared and is attached hereto.

In the left-hand column below is the column and line number where errors occurred in the patent. In the right-hand column is the page and line number in the application where the correct information appears.

Patent Reads:

Column 2, line 13:

“relate d to”

Column 2, line 18:

“even though has been”

Application Should Read:

Page 2, line 25:

--related to--

Page 3, line 4:

--even though it has been--

Column 5, line 45:

“results altered”

Column 10, line 17:

“derived form”

Column 13, lines 10-12:

“fatty oils, for example, sesame oil, or synthetic fatty acid esters, for example, sesame oil, or synthetic fatty acid esters, for example, ethyl”

Column 15, line 60:

“transferred it to”

Column 17, line 1:

“depends of the”

Column 17, line 9:

“adjusted to pH to 4.5”

Column 17, line 31:

“exchange is carried out”

Patent Reads:Column 19, line 26:

“was ravaged”

Patent Reads:Column 19, line 55:

“Paaavola”

Page 9, line 24:

--results are altered--

Page 18, line 24:

--derived from--

Page 24, lines 15-17:

--fatty oils, for example, sesame oil, or synthetic fatty acid esters, for example, ethyl--

Page 29, line 23:

--transferred to--

Page 32, line 2:

--depends on the--

Page 32, line 9:

--adjusted to pH 4.5--

Page 33, line 1:

--exchange and carried out--

Application Reads:Page 36, line 17:

--was lavaged--

Application Should Read:Page 37, line 10:

--Paavola--

Patent Reads:Column 20, line 33:“400 \parallel L”**Patent Reads:**Column 22, line 37:

“mutated in way that”

Patent Reads:Column 24, line 22:

“6569-4584”

Application Reads:Page 38, line 16:--400 μ L--**Application Should Read:**Page 42, line 10:

--mutated in a way that--

Application Reads:Page 46, line 20:

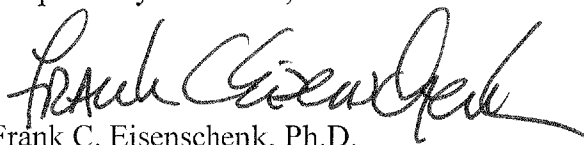
--6569-6584--.

A true and correct copy of pages 36, 38, and 46 of the specification as filed which support Applicants' assertion of the errors on the part of the Patent Office accompanies this Certificate of Correction.

The fee of \$100.00 was paid at the time this Request was filed. The Commissioner is also authorized to charge any additional fees as required under 37 CFR 1.20(a) to Deposit Account No. 19-0065.

Approval of the Certificate of Correction is respectfully requested.

Respectfully submitted,



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FCE/sl

Attachments: Copy of pages 36, 38, and 46 of the specification
Certificate of Correction

(saline). The mice were sacrificed on at 4 hours after the CCL2 or CCL2-P8A final injection.

The assay for thioglycollate-induced peritoneal cell recruitment has been published (Mishell B, 1980). Briefly, thioglycollate medium was prepared by suspending
5 30 g of dehydrated thioglycollate medium (Becton Dickinson) in 1 liter of cold distilled water, then heated until boiling to dissolve the powder completely. The medium was then aliquoted into 100 ml bottles and autoclaved. After cooling, the medium was stored in the dark at room temperature for at least one month. Cellular recruitment was induced by intraperitoneal injection of mice in groups of 3 with 200 μ l of a 3% solution
10 of thioglycollate on Day 1, 30 minutes after CCL2*-P8A administration. CCL2* was administered daily thereafter for 3 days (Days 2, 3 and 4). Dexamethasone (Sigma) was used as a positive control and administered at 10 mg/kg intraperitoneally. The mice were sacrificed on Day 5.

Peritoneal lavages to assess cell recruitment in the previous assays were
15 performed as follows. Mice were sacrificed by asphyxiation with rising concentrations of CO₂ in a plexiglass box. Skin was cleaned with 70% ethanol. The outer layer of skin was removed, exposing the peritoneal membrane. The peritoneal cavity was lavaged 3 times with 5 ml ice cold PBS (phosphate buffered saline) and fluid was pooled in a 15 ml polystyrene Falcon tube (Becton Dickinson) on ice. Each lavage was accompanied
20 with a light massage of the peritoneal cavity. Lavage fluid was centrifuged at 425xg, the supernatant discarded and the resultant cell pellet was resuspended by gentle multiple pipetting in 1 ml PBS. 10 μ l cell suspension was stained with 90 μ l trypan blue and total cell counts were enumerated with a Neubauer haemocytometer by counting 4 areas each of 1 mm². The mean of the 4 counts was taken, multiplied by the dilution factor of
25 10, and multiplied again by 10 to give the number of cells per μ l, according to the

total volume of 200 μ l, which were prepared by mixing 25 μ l ovalbumin (2mg/ml), 250 μ l aluminium hydroxide in 725 μ l LPS-free 0.9% NaCl (saline) and precipitated 3-4 hours at 4°C. Fifteen days after sensitisation, mice were treated and challenged in groups of 6 mice with the intranasal administration of 15 μ g ovalbumin in 50 μ l saline, under inhaled anaesthesia (Isoflurane) daily from day 15 to 19. CCL2-P8A (200 μ l, 10 μ g per intraperitoneal injection) was administered 30 minutes before each challenge. Peritoneal lavages to assess cell recruitment and cell counts were performed as described above in Example 2.

EAE (Experimental Autoimmune Encephalomyelitis) model

C57Bl/6 mice from Charles River Italy (the selected strain has documented susceptibility to EAE; Sahrbacher UC et al., 1998.) are immunized (day=0) by injecting s.c. in the left flank 0.2 mL of an emulsion composed of 200 μ g MOG₃₅₋₅₅ peptide (Neosystem, Strasbourg, France) in Complete Freund's Adjuvant (CFA, Difco, Detroit, U.S.A.) containing 0.5 mg of *Mycobacterium tuberculosis*. Immediately after, they receive an i.p. injection of 500 ng pertussis toxin (List Biological Lab., Campbell, CA, U.S.A.) dissolved in 400 μ L of buffer (0.5 M NaCl, 0.017% Triton X-100, 0.015 M Tris, pH=7.5). On day 2, the animals are given a second i.p. injection of 500 ng pertussis toxin. On day 7, the mice receive a second dose of 200 μ g of MOG₃₅₋₅₅ peptide in CFA injected s.c. in the right flank. Starting approximately from day 8-10, this procedure results in a gradually progressing paralysis, arising from the tail and ascending up to the forelimbs. Starting from day 7 the animals are individually examined for the presence of paralysis by means of a clinical score as follows:

0 = no sign of disease

0.5 = partial tail paralysis

1 = tail paralysis

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UNITED STATES PATENT AND TRADEMARK OFFICE

CERTIFICATE OF CORRECTION

PATENT NO. : 7,740,833

Page 1 of 2

APPLICATION NO.: 10/573,625

DATED : June 22, 2010

INVENTORS : Amanda Proudfoot, Jeffrey Shaw, Zoe Johnson

It is certified that errors appear in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

Column 2,

Line 13, "relate d to" should read --related to--.

Line 18, "even though has been" should read --even though it has been--.

Column 5,

Line 45, "results altered" should read --results are altered--.

Column 10,

Line 17, "derived form" should read --derived from--.

Column 13,

Lines 10-12, "fatty oils, for example, sesame oil, or synthetic fatty acid esters, for example, sesame oil, or synthetic fatty acid esters, for example, ethyl" should read --fatty oils, for example, sesame oil, or synthetic fatty acid esters, for example, ethyl--.

Column 15,

Line 60, "transferred it to" should read --transferred to--.

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Page 2 of 2

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Column 17,

Line 1, "depends of the" should read --depends on the--.

Line 9, "adjusted to pH to 4.5" should read --adjusted to pH 4.5--.

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Line 55, "Paaavola" should read --Paavola--.

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Column 24,

Line 22, "6569-4584" should read --6569-6584--.

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